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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/532,831	03/09/2006	Hans-Ulrich Petereit	267336US0PCT	8866
OBLON, SPIVAK, MCCLELLAND MAIER & NEUSTADT, L.L.P. 1940 DUKE STREET			EXAMINER	
			WESTERBERG, NISSA M	
ALEXANDRIA, VA 22314			ART UNIT	PAPER NUMBER
		1618		
			NOTIFICATION DATE	DELIVERY MODE
			02/22/2010	ELECTRONIC

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

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Advisory Action Before the Filing of an Appeal Brief

Application No.	Applicant(s)		
10/532,831	PETEREIT ET AL.		
Examiner	Art Unit		
I	1	l	

	Missa W. Westerberg	1010	
The MAILING DATE of this communication app	ears on the cover sheet with the	correspondence add	ress
THE REPLY FILED <u>03 February 2010</u> FAILS TO PLACE THIS	APPLICATION IN CONDITION FO	R ALLOWANCE.	
1. The reply was filed after a final rejection, but prior to or o application, applicant must timely file one of the following application in condition for allowance; (2) a Notice of App for Continued Examination (RCE) in compliance with 37 periods:	replies: (1) an amendment, affidavi eal (with appeal fee) in compliance	t, or other evidence, wwith 37 CFR 41.31; or	which places the r (3) a Request
a) The period for reply expiresmonths from the mailir	ng date of the final rejection.		
b) The period for reply expires on: (1) the mailing date of this no event, however, will the statutory period for reply expire Examiner Note: If box 1 is checked, check either box (a) or MONTHS OF THE FINAL REJECTION. See MPEP 706.07	later than SIX MONTHS from the mailing (b). ONLY CHECK BOX (b) WHEN THE (f).	g date of the final rejection FIRST REPLY WAS FI	on. LED WITHIN TWO
Extensions of time may be obtained under 37 CFR 1.136(a). The date have been filed is the date for purposes of determining the period of extender 37 CFR 1.17(a) is calculated from: (1) the expiration date of the set forth in (b) above, if checked. Any reply received by the Office late may reduce any earned patent term adjustment. See 37 CFR 1.704(b) NOTICE OF APPEAL	xtension and the corresponding amount shortened statutory period for reply origing than three months after the mailing dat	of the fee. The appropria nally set in the final Offic	ate extension fee be action; or (2) as
2. The Notice of Appeal was filed on A brief in com	pliance with 37 CFR 41.37 must be	filed within two months	s of the date of
filing the Notice of Appeal (37 CFR 41.37(a)), or any extension Notice of Appeal has been filed, any reply must be filed to AMENDMENTS			e appeal. Since a
3. The proposed amendment(s) filed after a final rejection,			cause
(a) They raise new issues that would require further co	•	ΓE below);	
 (b) ☐ They raise the issue of new matter (see NOTE belet) (c) ☐ They are not deemed to place the application in beappeal; and/or 	•	ducing or simplifying t	he issues for
(d) ☐ They present additional claims without canceling a NOTE: (See 37 CFR 1.116 and 41.33(a))		ected claims.	
4. The amendments are not in compliance with 37 CFR 1.		mpliant Amendment (PTOL-324).
Applicant's reply has overcome the following rejection(s):		
6. Newly proposed or amended claim(s) would be a non-allowable claim(s).			_
7. For purposes of appeal, the proposed amendment(s): a) how the new or amended claims would be rejected is pro The status of the claim(s) is (or will be) as follows: Claim(s) allowed:		I be entered and an e	xpianation of
Claim(s) objected to:			
Claim(s) rejected: Claim(s) withdrawn from consideration:			
AFFIDAVIT OR OTHER EVIDENCE			
 The affidavit or other evidence filed after a final action, b because applicant failed to provide a showing of good ar was not earlier presented. See 37 CFR 1.116(e). 			
9. The affidavit or other evidence filed after the date of filing entered because the affidavit or other evidence failed to showing a good and sufficient reasons why it is necessa	overcome <u>all</u> rejections under appea	al and/or appellant fail	s to provide a
10. ☐ The affidavit or other evidence is entered. An explanation REQUEST FOR RECONSIDERATION/OTHER	on of the status of the claims after e	ntry is below or attach	ed.
11. The request for reconsideration has been considered b See Continuation Sheet.	ut does NOT place the application ir	condition for allowan	ce because:
12. ☐ Note the attached Information <i>Disclosure Statement</i>(s).13. ☐ Other:	(PTO/SB/08) Paper No(s)		
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	/Jake M. Vu/ Primary Examiner, Art U	Init 1618	

Continuation of 11. does NOT place the application in condition for allowance because:

Claims 1 - 3, 8, 9, 13, 14 and 16 are rejected under 35 USC 103(a) as being unpatentable over Ulmius et al. (US 5,643,602) in view of Beckert et al. (WO 01/68058). Applicants argue there is no direction to select the claimed metarial, contrary to the assertion of the previous Office Action there is sufficient evidence of unexpected results, there is no reasonable prediction of success for the percentage release of active agent and there is no support for the exclusion of the various excipients.

These arguments are unpersuasive. Ulmius et al. includes the EUDRAGIT® NE in aquesous diseprsion (what the '30D' part of the name refers to) among a limited number of preferred polymers and based on Applicant's arguments, it appears that this polymer is the ingredient which provides the independence of the release rate from the osmotic/ionic conditions of the medium. The teachings of the reference aren't limited to the examples. The exmamples given by Applicant to support the unexpected percentage release rate all relate to a very limited number of active ingredients (all but one relate to budesonide) and there is insiufficient evidence that the observed independence of release rate from the osmotic conditions applies to all active ingredients, which the breadth of the claims in regards to the active ingredient. The perecentage release flows from the comopsition and structure of the dosage form, and there is a reasonable expectation of success in preparing the dosage form, which will lead to the release profile. In regards to the excipients, Ulmius discloses that they are optional, which means that these ingredients are not required and thus can be excluded from the composition(s).

Claims 1 - 3, 8, 9, 13, 14 and 16 are rejected nder 35 USC 103(a) as being unpatentable over Ulmius et al. (US 5,643,602) in view of Gang et al. (Proc of the 7th SECJ, 2001). Applicants response is slightly confusing as to this rejection as a p 7 of the response indicates that Beckert is the primary reference, but then references Ulmius on p 8. As no rejection with Beckert as the primary refrence has been made, the Examiner will assume that the primary reference being argued was Ulmius.

Applicants argue that the active in Gang is in the pellet core and not bound to the inner coating layer, which leads to different release curves and again that the observed relese profile is unexpected.

These arguments are unpersuasive. The location of the active ingredient comes from the teachings of the primary reference (Ulmius). The arguments regarding the unexpected release profile were discussed above.